

K960963 CORD WRAP WITH NEEDLE RESHEATHERMay 6, 1996
56 days to decisionK960963 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k960963/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Mar 11, 1996
Decision date	May 6, 1996
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Osborn Medical Corp.
Location	Utica, MN, US
Contact	BILL D DAVIS
510(k) history	2 submissions · 2 cleared · 1996-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k960963/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 3, 2026